12 December 2023
EMA/515639/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): clarithromycin

Procedure No. PSUSA/00000788/202304

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Clarithromycin 1A Pharma 500 mg - Filmtabletten	not available	1-26245	1A PHARMA GMBH	AT
Klacid - Lactobionat 0,5 g - Trockensubstanz zur Infusionsbereitung	not available	1-20013	MYLAN ÖSTERREICH GMBH	AT
Clarithromycin Sandoz 250 mg/5 ml – Granulat für orale Suspension	NL/H/2144/002	1-25596	SANDOZ GMBH	AT
Clarithromycin Sandoz 125 mg/5 ml – Granulat für orale Suspension	NL/H/2144/001	1-25595	SANDOZ GMBH	AT
Klacid 125 mg / 5 ml - Granulat für orale Suspension	not available	1-19911	MYLAN ÖSTERREICH GMBH	AT
Klacid 250 mg / 5 ml - Granulat für orale Suspension	not available	1-21047	MYLAN ÖSTERREICH GMBH	AT
Klacid Uno – Filmtabletten mit veränderter Wirkstoff-Freisetzung	IE/H/0105/001	1-22085	MYLAN ÖSTERREICH GMBH	AT
Clarithromycin Sandoz 500 mg - Filmtabletten	not available	1-26244	SANDOZ GMBH	AT
Klacid 250 mg - Filmtabletten	not available	1-19163	MYLAN ÖSTERREICH GMBH	AT
KLACID 500 mg - Filmtabletten	not available	1-21527	MYLAN ÖSTERREICH GMBH	AT
Maclar 500 mg, Filmtabletten	not available	BE173467	MYLAN EPD BVBA/SPRL	BE
Maclar 500 mg, filmomhulde tabletten	not available	BE173467	MYLAN EPD BVBA/SPRL	BE
Maclar 500 mg, comprimés pelliculés	not available	BE173467	MYLAN EPD BVBA/SPRL	BE
Heliclar 500 mg, Filmtabletten	not available	BE173451	MYLAN EPD BVBA/SPRL	BE
Heliclar 500 mg, filmomhulde tabletten	not available	BE173451	VIATRIS HEALTHCARE	BE
Heliclar 500 mg, comprimés pelliculés	not available	BE173451	MYLAN EPD BVBA/SPRL	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Monoclarium 200 mg, gélule à libération prolongée.	BE/H/0365/001	BE312417	LABORATOIRES SMB S.A.	BE
Monoclarium 200 mg, harde capsules met verlengde afgifte.	BE/H/0365/001	BE312417	LABORATOIRES SMB S.A.	BE
Monoclarium 200 mg Hartkapsel, retardiert	BE/H/0365/001	BE312417	LABORATOIRES SMB S.A.	BE
Biclar 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	BE165575	VIATRIS HEALTHCARE	BE
Biclar Kids 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	BE215092	VIATRIS HEALTHCARE	BE
Biclar 125 mg/5 ml, granulaat voor orale suspensie	not available	BE165575	VIATRIS HEALTHCARE	BE
Biclar Kids 250 mg/5ml, granulaat voor orale suspensie	not available	BE215092	VIATRIS HEALTHCARE	BE
Biclar 125 mg/5 ml, granulés pour suspension buvable	not available	BE165575	VIATRIS HEALTHCARE	BE
Biclar Kids 250 mg/5ml, granulés pour suspension buvable	not available	BE215092	VIATRIS HEALTHCARE	BE
Biclar I.V. 500 mg, Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	not available	BE173476	VIATRIS HEALTHCARE	BE
Biclar I.V. 500 mg, poeder voor concentraat voor oplossing voor infusie	not available	BE173476	VIATRIS HEALTHCARE	BE
Biclar I.V. 500 mg, poudre pour solution à diluer pour perfusion	not available	BE173476	VIATRIS HEALTHCARE	BE
Biclar Uno 500 mg Tabletten mit veränderter Wirkstofffreisetzung	IE/H/0105/001	BE187144	VIATRIS HEALTHCARE	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Biclar Uno 500 mg, comprimés à libération modifiée	IE/H/0105/001	BE187144	VIATRIS HEALTHCARE	BE
Biclar Uno 500 mg tabletten met gereguleerde afgifte	IE/H/0105/001	BE187144	VIATRIS HEALTHCARE	BE
Biclar 250 mg, Filmtabletten	not available	BE154086	VIATRIS HEALTHCARE	BE
Biclar Forte 500 mg, Filmtabletten	not available	BE173485	VIATRIS HEALTHCARE	BE
Biclar 250 mg, filmomhulde tabletten	not available	BE154086	VIATRIS HEALTHCARE	BE
Biclar Forte 500 mg, filmomhulde tabletten	not available	BE173485	VIATRIS HEALTHCARE	BE
Biclar 250 mg, comprimés pelliculés	not available	BE154086	VIATRIS HEALTHCARE	BE
Biclar Forte 500 mg, comprimés pelliculés	not available	BE173485	VIATRIS HEALTHCARE	BE
Lekoklar 500 mg film-coated tablets	NL/H/2087/002	20110649	SANDOZ PHARMACEUTICALS D.D.	BG
Клацид 125 mg/5 ml гранули за перорална суспензия	not available	20010617	MYLAN EOOD	BG
Клацид 250 mg/5 ml гранули за перорална суспензия	not available	20050512	MYLAN EOOD	BG
Клацид 250 mg филмирани таблетки	not available	20010734	MYLAN EOOD	BG
Клацид SR 500 mg таблетки с изменено освобождаване	not available	20010517	MYLAN EOOD	BG
Klacid 125 mg/5 ml granule pro perorální suspenzi	not available	15/634/94-C	VIATRIS HEALTHCARE LTD	CZ
Klacid 250 mg/5 ml granule pro perorální suspenzi	not available	15/355/98-C	VIATRIS HEALTHCARE LTD	CZ
Klacid 500 mg prášek pro koncentrát pro infuzní roztok	not available	15/236/99-C	VIATRIS HEALTHCARE LTD	CZ
Klacid 250 mg potahované tablety	not available	15/1157/94-C	VIATRIS HEALTHCARE LTD	CZ
Klacid 500 mg potahované tablety	not available	15/374/97-C	VIATRIS HEALTHCARE LTD	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid SR 500 mg tablety s řízeným uvolňováním	not available	15/062/98-C	VIATRIS HEALTHCARE LTD	CZ
Klacid 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	not available	46110.00.00	VIATRIS HEALTHCARE GMBH	DE
Klacid Saft, 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	19219.00.01	VIATRIS HEALTHCARE GMBH	DE
Klacid Saft Forte 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	40373.00.00	VIATRIS HEALTHCARE GMBH	DE
Klacid 500 mg Pulver und Lösungsmittel für ein Konzentrat zur Herstellung einer Infusionslösung	not available	93335.00.00	VIATRIS HEALTHCARE GMBH	DE
Klacid Pro, 250 mg Filmtabletten	not available	28401.00.00	VIATRIS HEALTHCARE GMBH	DE
Klacid Filmtabletten, 250 mg	not available	19219.00.00	VIATRIS HEALTHCARE GMBH	DE
Klacid, filmovertrukne tabletter	not available	15527	VIATRIS APS	DK
Klacid, granulat til oral suspension	not available	18018	VIATRIS APS	DK
Klacid, granulat til oral suspension	not available	15382	VIATRIS APS	DK
Klacid Uno, depottabletter	IE/H/0105/001	18861	VIATRIS APS	DK
Klacid, pulver til infusionsvæske, opløsning	not available	30742	VIATRIS APS	DK
Klacid 125 mg/5 ml, suukaudse suspensiooni graanulid	not available	268799	MYLAN HEALTHCARE SIA	EE
Klacid i.v., 500 mg infusioonilahuse kontsentraadi pulber	not available	312800	MYLAN HEALTHCARE SIA	EE
Klacid 250 mg, õhukese polümeerikattega tabletid	not available	268699	MYLAN HEALTHCARE SIA	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid SR 500 mg, toimeainet modifitseeritult vabastavad tabletid	not available	268999	MYLAN HEALTHCARE SIA	EE
Klacid, 500 mg õhukese polümeerikattega tabletid	not available	751011	MYLAN HEALTHCARE SIA	EE
Klacid 500 mg liofilizado para solución para perfusión	not available	60202	MYLAN IRE HEALTHCARE LIMITED	ES
Klacid 500 mg granulado para suspensión oral	not available	60518	MYLAN IRE HEALTHCARE LIMITED	ES
Klacid 500 mg comprimidos recubiertos con película	not available	60515	MYLAN IRE HEALTHCARE LIMITED	ES
Klacid unidía 500 mg comprimidos de liberación modificada	not available	63380	MYLAN IRE HEALTHCARE LIMITED	ES
Klacid 250 mg tabletti, kalvopäällysteinen	not available	11451	VIATRIS OY	FI
Klacid 500 mg tabletti, kalvopäällysteinen	not available	11923	VIATRIS OY	FI
Klacid 500 mg filmdragerade tabletter	not available	11923	VIATRIS OY	FI
Klacid 250 mg filmdragerade tabletter	not available	11451	VIATRIS OY	FI
Zeclar OD 500 mg tabletter med modifierad frisättning	not available	12844	ORION CORPORATION	FI
Zeclar OD 500 mg säädellysti vapauttavat tabletit	not available	12844	ORION CORPORATION	FI
Zeclar 250 mg filmdragerade tabletter	not available	12341	ORION CORPORATION	FI
Zeclar 500 mg filmdragerade tabletter	not available	12342	ORION CORPORATION	FI
Zeclar 50 mg/ml granulat till oral suspension	not available	13090	ORION CORPORATION	FI
Zeclar 250 mg kalvopäällysteiset tabletit	not available	12341	ORION CORPORATION	FI
Zeclar 500 mg kalvopäällysteiset tabletit	not available	12342	ORION CORPORATION	FI
Zeclar 50 mg/ml rakeet oraalisuspensiota varten	not available	13090	ORION CORPORATION	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid 50 mg/ml rakeet oraalisuspensiota varten	not available	13089	VIATRIS OY	FI
Klacid 50 mg/ml granulat till oral suspension	not available	13089	VIATRIS OY	FI
Klacid OD 500 mg säädellysti vapauttava tabletti	IE/H/0105/001	12843	VIATRIS OY	FI
KLACID OD 500 mg tablett med modifierad frisättning	IE/H/0105/001	12843	VIATRIS OY	FI
Klacid 500 mg kuiva-aine välikonsentraatiksi infuusionestettä varten, liuos	not available	14371	VIATRIS OY	FI
Klacid 500 mg pulver till koncentrat till infusionsvätska, lösning	not available	14371	VIATRIS OY	FI
ZECLAR 0,5 g, poudre pour solution à diluer pour perfusion	not available	34009 355 843 5 9	VIATRIS MEDICAL	FR
ZECLAR 0,5 g, poudre pour solution à diluer pour perfusion	not available	34009 346 439 0 3	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 564 0 6	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 347 830 5 0	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 340 974 1 6	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 347 831 1 1	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 565 7 4	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 566 3 5	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 568 6 4	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34008 340 975 8 4	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 569 2 5	VIATRIS MEDICAL	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 570 0 7	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 571 7 5	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 572 3 6	VIATRIS MEDICAL	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 348 736 2 1	VIATRIS MEDICAL	FR
ZECLAR 25 mg/ml, granulés pour suspension buvable	not available	34009 345653 9 7	VIATRIS MEDICAL	FR
ZECLAR 50 mg/ml, granulés pour suspension buvable	not available	34009 347 855 8 0	VIATRIS MEDICAL	FR
ZECLAR 50 mg/ml, granulés pour suspension buvable	not available	34009 347 853 5 1	VIATRIS MEDICAL	FR
ZECLAR 25 mg/ml, granulés pour suspension buvable	not available	34009 345655 1 9	VIATRIS MEDICAL	FR
CLARITHROMYCINE ABBOTT 25 mg/ml, granulés pour suspension buvable	not available	NL22348	ABBOTT BIOLOGICALS B.V. (WEESP)	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 533 9 2	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 565 957 7 8	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 565 958 3 9	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 539 7 2	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 537 4 3	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 534 5 3	VIATRIS MEDICAL	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 535 1 4	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 536 8 2	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 541 1 5	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 540 5 4	VIATRIS MEDICAL	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 538 0 4	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 346 497 0 7	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 280 6 6	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 281 2 7	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 013 8 0	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 303 6 6	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 207 3 5	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 346 498 7 5	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 205 0 6	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 279 8 4	VIATRIS MEDICAL	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 344 844 5 2	VIATRIS MEDICAL	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 344 845 1 3	VIATRIS MEDICAL	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZECLAR 250 mg, comprimé pelliculé	not available	34009 334 031 1 9	VIATRIS MEDICAL	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 344 846 8 1	VIATRIS MEDICAL	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 344 847 4 2	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 204 4 5	VIATRIS MEDICAL	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 206 7 4	VIATRIS MEDICAL	FR
KLARICID® 500 mg/vial Κόνις για πυκνό σκεύασμα για παρασκευή διαλύματος προς έγχυση	not available	36469/10/18-03-2011	VIATRIS HEALTHCARE LTD	GR
BREVIL® 200 mg ?a????a pa?atetaμ???? ap?d?sμe?s??, s?????	FI/H/0624/001	62178/14/09/2010	MEDITRINA PHARMACEUTICALS LTD	GR
KLARICID®250mg/5ml Κοκκία για πόσιμο εναιώρημα	not available	36464/10/18-03-2011	VIATRIS HEALTHCARE LTD	GR
KLARICID® OD 500 mg Δισκία Ελεγχόμενης Αποδέσμευσης	not available	36472/10/18-03-2011	VIATRIS HEALTHCARE LTD	GR
KLARICID® 500 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	36466/10/18-03-2011	VIATRIS HEALTHCARE LTD	GR
KLARICID® 250 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	36468/10/18-03-2011	VIATRIS HEALTHCARE LTD	GR
MAKCIN 500 mg filmom obložene tablete	not available	HR-H-696573111	BELUPO D.D.	HR
MAKCIN SR 500 mg tablete s produljenim oslobađanjem	not available	HR-H-030175902	BELUPO D.D.	HR
Klacid 250 mg filmtabletta	not available	OGYI-T-2200/05	VIATRIS HEALTHCARE LTD	HU
Klacid 250 mg filmtabletta	not available	OGYI-T-2200/04	VIATRIS HEALTHCARE LTD	HU
Klacid 500 mg filmtabletta	not available	OGYI-T-2200/06	VIATRIS HEALTHCARE LTD	HU
Klacid UNO 500 mg retard filmtabletta	not available	OGYI-T-2200/08	VIATRIS HEALTHCARE LTD	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid 125 mg/5 ml granulátum 100 ml belsőleges szuszpenzióhoz	not available	OGYI-T-2200/02	VIATRIS HEALTHCARE LTD	HU
Klacid Baby 125 mg/5 ml granulátum 60 ml belsőleges szuszpenzióhoz	not available	OGYI-T-2200/01	VIATRIS HEALTHCARE LTD	HU
Klacid Kid 250 mg/5 ml granulátum 70 ml belsőleges szuszpenzióhoz	not available	OGYI-T-2200/03	VIATRIS HEALTHCARE LTD	HU
Klacid 50 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-2200/10	VIATRIS HEALTHCARE LTD	HU
Klacid IV 500mg Powder for Concentrate for Solution for Infusion	not available	PA 2010/4/3	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid LA 500mg Modified release tablets	IE/H/0105/001	PA 2010/4/4	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid Paediatric Suspension 125mg/5ml, Granules for Oral Suspension	not available	PA 2010/4/5	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid 250mg/5ml Granules for Oral Suspension	not available	PA 2010/4/6	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid 250 mg Film-coated Tablets	not available	PA 2010/4/1	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid Forte 500mg Film-coated Tablets	not available	PA 2010/4/2	MYLAN IRE HEALTHCARE LIMITED	IE
MACLADIN RM 500 mg Compresse a rilascio modificato	not available	027530144	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 500 mg Granulato per sospensione orale	not available	027530106	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 250 mg Compresse rivestite	not available	027530056	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 500 mg Compresse rivestite	not available	027530118	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 250 mg Granulato per sospensione orale	not available	027530094	LABORATORI GUIDOTTI S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MACLADIN 125 mg/5 ml Granulato per sospensione orale	not available	027530068	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 250 mg/5 ml Granulato per sospensione orale	not available	027530120	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 500 mg/10 ml Polvere e solvente per concentrato per soluzione per infusione	not available	027530043	LABORATORI GUIDOTTI S.P.A.	IT
KLACID 125 mg/5 ml Granulato per sospensione orale	not available	027370067	MYLAN ITALIA S.R.L.	IT
KLACID 250 mg/5 ml Granulato per sospensione orale	not available	027370117	MYLAN ITALIA S.R.L.	IT
KLACID 250 mg Granulato per sospensione orale	not available	027370093	MYLAN ITALIA S.R.L.	IT
KLACID 500 mg Granulato per sospensione orale	not available	027370105	MYLAN ITALIA S.R.L.	IT
SORICLAR 250 mg Compresse rivestite	not available	037456050	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 500 mg Compresse rivestite	not available	037456062	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 125 mg/5 ml Granulato per sospensione orale	not available	037456011	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg/5 ml Granulato per sospensione orale	not available	037456023	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg Granulato per sospensione orale	not available	037456086	ABIOGEN PHARMA S.P.A.	IT
SORICLAR RM 500 mg Compresse a rilascio modificato	not available	037456098	ABIOGEN PHARMA S.P.A.	IT
VECLAM RM 500 mg Compresse a rilascio modificato	not available	027529130	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VECLAM 500 mg Compresse rivestite	not available	027529116	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 250 mg Compresse rivestite	not available	027529054	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 500 mg Granulato per sospensione orale	not available	027529092	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 250 mg Granulato per sospensione orale	not available	027529080	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 125 mg/5 ml Granulato per sospensione orale	not available	027529041	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 250 mg/5 ml Granulato per sospensione orale	not available	027529104	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 500 mg/10 ml Polvere e solvente per concentrato per soluzione per infusione	not available	027529039	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
KLACID 250 mg Compresse rivestite	not available	027370055	MYLAN ITALIA S.R.L.	IT
KLACID 500 mg Compresse rivestite	not available	027370129	MYLAN ITALIA S.R.L.	IT
KLACID RM 500 mg Compresse a rilascio modificato	not available	027370143	MYLAN ITALIA S.R.L.	IT
KLACID 500 mg/10 ml Polvere e solvente per soluzione per infusione	not available	027370042	MYLAN ITALIA S.R.L.	IT
WINCLAR 500 mg Compresse rivestite	not available	037446061	I.B.N. SAVIO S.R.L.	IT
WINCLAR 250 mg/5 mL granulato per sospensione orale	not available	037446022	I.B.N. SAVIO S.R.L.	IT
Klacid i.v. 500 mg milteliai infuziniam tirpalui	not available	LT/1/98/0264/007	MYLAN HEALTHCARE SIA	LT
Klacid 125 mg/5 ml granulės geriamajai suspensijai	not available	LT/1/98/0264/005	MYLAN HEALTHCARE SIA	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid 125 mg/5 ml granulės geriamajai suspensijai	not available	LT/1/98/0264/004	MYLAN HEALTHCARE SIA	LT
Klacid 250 mg plėvele dengtos tabletės	not available	LT/1/98/0264/006	MYLAN HEALTHCARE SIA	LT
Klacid SR 500 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/98/0264/002	MYLAN HEALTHCARE SIA	LT
Klacid SR 500 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/98/0264/003	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/009	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/010	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/011	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/012	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/013	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/014	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/015	MYLAN HEALTHCARE SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/008	MYLAN HEALTHCARE SIA	LT
Maclar 500 mg, comprimés pelliculés	not available	1996060327	VIATRIS HEALTHCARE	LU
Maclar 500 mg, Filmtabletten	not available	1996060327	VIATRIS HEALTHCARE	LU
Monoclarium 200 mg, gélule à libération prolongée	BE/H/0365/001	2008110062	LABORATOIRES SMB S.A.	LU
Monoclarium 200 mg Hartkapsel, retardiert	BE/H/0365/001	2008110062	LABORATOIRES SMB S.A.	LU
Biclar I.V. 500 mg, poudre pour solution à diluer pour perfusion	not available	1996060326	VIATRIS HEALTHCARE	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Biclar I.V. 500 mg, Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	not available	1996060326	VIATRIS HEALTHCARE	LU
Biclar 125 mg/5 ml, granulés pour suspension buvable	not available	1994110818	VIATRIS HEALTHCARE	LU
Biclar Kids 250 mg/5ml, granulés pour suspension buvable	not available	2001010001	VIATRIS HEALTHCARE	LU
Biclar 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	1994110818	VIATRIS HEALTHCARE	LU
Biclar Kids 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	2001010001	VIATRIS HEALTHCARE	LU
Heliclar 500 mg, comprimés pelliculés	not available	1996060325	VIATRIS HEALTHCARE	LU
Heliclar 500 mg, Filmtabletten	not available	1996060325	VIATRIS HEALTHCARE	LU
Biclar Uno 500 mg, comprimés à libération modifiée	IE/H/0105/001	2005106594	VIATRIS HEALTHCARE	LU
Biclar Uno 500 mg Tabletten mit veränderter Wirkstofffreisetzung Clarithromycin	IE/H/0105/001	2005106594	VIATRIS HEALTHCARE	LU
Biclar 250 mg, comprimés pelliculés	not available	1997124275	VIATRIS HEALTHCARE	LU
Biclar Forte 500 mg, comprimés pelliculés	not available	1996060324	VIATRIS HEALTHCARE	LU
Biclar 250 mg, Filmtabletten	not available	1997124275	VIATRIS HEALTHCARE	LU
Biclar Forte 500 mg, Filmtabletten	not available	1996060324	VIATRIS HEALTHCARE	LU
Klacid i.v. 500 mg pulveris infūziju šķīduma koncentrāta pagatavošanai	not available	94-0306	MYLAN HEALTHCARE SIA	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid 250 mg apvalkotās tabletes	not available	94-0307	MYLAN HEALTHCARE SIA	LV
Klacid SR 500 mg ilgstošās darbības tabletes	not available	98-0691	MYLAN HEALTHCARE SIA	LV
Klacid 500 mg apvalkotās tabletes	not available	10-0609	MYLAN HEALTHCARE SIA	LV
Klacid 125 mg/5 ml granulas iekšķīgi lietojamas suspensijas pagatavošanai	not available	94-0308	MYLAN HEALTHCARE SIA	LV
KLACID XL 500 mg Modified-release tablets	not available	MA1507/01305	VIATRIS HEALTHCARE LTD	MT
KLACID 500 mg Coated Tablets	not available	MA1507/01302	VIATRIS HEALTHCARE LTD	MT
KLACID 250 mg Coated tablets	not available	MA1507/01301	VIATRIS HEALTHCARE LTD	MT
Klacid IV 500mg Powder for Concentrate for Solution for Infusion	not available	MA1507/01306	VIATRIS HEALTHCARE LTD	MT
Klacid Paediatric Suspension 125mg/5ml, Granules for Oral Suspension	not available	MA1507/01303	VIATRIS HEALTHCARE LTD	MT
Klacid 250mg/5ml Granules for Oral Suspension	not available	MA1507/01304	VIATRIS HEALTHCARE LTD	MT
Klacid SR, 500 mg tablet met gereguleerde afgifte	IE/H/0105/001	RVG 21555	MYLAN HEALTHCARE B.V.	NL
Klacid 25 mg/ml granulat til mikstur	not available	94-1401	VIATRIS AS	NO
Klacid 50 mg/ml granulat til mikstur	not available	97-01645	VIATRIS AS	NO
Klacid OD 500 mg depottabletter	not available	97-2350	VIATRIS AS	NO
Klacid, 250 mg, tabletki powlekane	not available	R/3318	MYLAN HEALTHCARE SP. Z.O.O.	PL
Klacid Uno, 500 mg, tabletki o zmodyfikowanym uwalnianiu	not available	7930	MYLAN HEALTHCARE SP. Z.O.O.	PL
Klacid, 500 mg, tabletki powlekane	not available	R/7194	MYLAN HEALTHCARE SP. Z.O.O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid, 125 mg/5 ml, granulat do sporządzania zawiesiny doustnej	not available	R/3317	MYLAN HEALTHCARE SP. Z.O.O.	PL
Klacid, 250 mg/5 ml, granulat do sporządzania zawiesiny doustnej	not available	7806	MYLAN HEALTHCARE SP. Z.O.O.	PL
Klacid, 500 mg, proszek do sporządzania koncentratu roztworu do infuzji	not available	R/3757	MYLAN HEALTHCARE SP. Z.O.O.	PL
Klacid Pediátrico 25 mg/ml granulado para suspensão oral	not available	2146983	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid Pediátrico 50 mg/ml granulado para suspensão oral	not available	2525384	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5248612	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5248620	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250204	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250212	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250220	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250238	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250246	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250253	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258504	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258470	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258462	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258454	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258447	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258439	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258421	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258413	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Klacid 250 mg comprimido revestido	not available	5546684	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid 500 mg comprimido revestido	not available	5546783	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3518784	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519386	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055687	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055182	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055281	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519287	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519188	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627586	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055083	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4056289	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4056081	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4056180	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055380	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519485	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4054987	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519584	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055984	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055885	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3518883	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3518982	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627487	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519089	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627685	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519683	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519782	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519881	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055489	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627784	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055588	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055786	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid 250 mg comprimate filmate	not available	8383/2015/01	VIATRIS HEALTHCARE LTD	RO
Klacid 250 mg comprimate filmate	not available	8383/2015/02	VIATRIS HEALTHCARE LTD	RO
Klacid SR 500 mg comprimate cu eliberare prelungită	not available	8384/2015/01	VIATRIS HEALTHCARE LTD	RO
Klacid SR 500 mg comprimate cu eliberare prelungită	not available	8384/2015/02	VIATRIS HEALTHCARE LTD	RO
Klacid SR 500 mg comprimate cu eliberare prelungită	not available	8384/2015/03	VIATRIS HEALTHCARE LTD	RO
Klacid I.V.500 mg pulbere pentru concentrat pentru soluţie perfuzabilă	not available	8385/2015/01	VIATRIS HEALTHCARE LTD	RO
Klacid 125 mg/5 ml granule pentru suspensie orală	not available	8386/2015/01	VIATRIS HEALTHCARE LTD	RO
Klacid 125 mg/5 ml granule pentru suspensie orală	not available	8386/2015/02	VIATRIS HEALTHCARE LTD	RO
Klacid 250 mg tabletter	not available	11372	BGP PRODUCTS AB	SE
Klacid 500 mg tabletter	not available	12164	BGP PRODUCTS AB	SE
KLACID I.V. 500 mg prášok na infúzny roztok	not available	15/0010/99-S	VIATRIS HEALTHCARE LTD	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
KLACID 125 mg/5 ml granulát na perorálnu suspenziu	not available	15/0283/97-S	VIATRIS HEALTHCARE LTD	SK
KLACID 250 mg/5 ml granulát na perorálnu suspenziu	not available	15/0179/99-S	VIATRIS HEALTHCARE LTD	SK
KLACID SR 500 mg tablety s riadeným uvoľňovaním	not available	15/0141/99-S	VIATRIS HEALTHCARE LTD	SK
KLACID 500 mg filmom obalené tablety	not available	15/0215/00-S	VIATRIS HEALTHCARE LTD	SK
Klaricid 250 mg Tablets	not available	PL 46302/0016	MYLAN PRODUCTS LIMITED	XI
Klaricid XL 500 mg Tablets	not available	PL 46302/0020	MYLAN PRODUCTS LIMITED	XI
Klaricid 500 mg Tablets	not available	PL 46302/0017	MYLAN PRODUCTS LIMITED	XI
Clarithromycin 250 mg/5ml suspension	NL/H/2144/002	PL 04416/0610	SANDOZ LTD	XI
Clarithromycin 125 mg/5ml suspension	NL/H/2144/001	PL 04416/0609	SANDOZ LTD	XI
Clarithromycin 250mg Film-coated Tablets	PT/H/2323/001	PL 04416/0641	SANDOZ LTD	XI
Klaricid IV 500 mg, Powder for Concentrate for Solution for Infusion	not available	PL 46302/0018	MYLAN PRODUCTS LIMITED	XI
Klaricid Paediatric Suspension 125mg/5ml	not available	PL 46302/0014	MYLAN PRODUCTS LIMITED	XI
Klaricid Paediatric Suspension 250mg/5ml	not available	PL 46302/0019	MYLAN PRODUCTS LIMITED	XI
Klaricid Adult Sachet 250mg	not available	PL 46302/0015	MYLAN PRODUCTS LIMITED	XI